

Durham Research Online

Deposited in DRO:

14 May 2020

Version of attached file:

Accepted Version

Peer-review status of attached file:

Peer-reviewed

Citation for published item:

Gregory, Joe and Verlet, Jan R. R. and Bull, James N. (2020) 'Spectroscopic characterisation of radical polyinterhalogen molecules.', *Physical chemistry chemical physics.*, 22 (16). pp. 8284-8288.

Further information on publisher's website:

<https://doi.org/10.1039/D0CP01311K>

Publisher's copyright statement:

Additional information:

Use policy

The full-text may be used and/or reproduced, and given to third parties in any format or medium, without prior permission or charge, for personal research or study, educational, or not-for-profit purposes provided that:

- a full bibliographic reference is made to the original source
- a [link](#) is made to the metadata record in DRO
- the full-text is not changed in any way

The full-text must not be sold in any format or medium without the formal permission of the copyright holders.

Please consult the [full DRO policy](#) for further details.

Cite this: DOI: 00.0000/xxxxxxxxxx

Spectroscopic characterisation of radical polyinterhalogen molecules[†]

Joe Gregory,^{*a} Jan R. R. Verlet^b and James N. Bull^a

Received Date

Accepted Date

DOI: 00.0000/xxxxxxxxxx

Spectroscopic characterisations of the radical polyinterhalogen molecules IF_2^- and I_2F^- are reported using anion photoelectron spectroscopy. The corresponding parent anions, IF_2^- and I_2F^- , are common products formed in hard Ar- CF_3I plasmas and are relevant in the semiconductor manufacture industry. The I_2F^- species, which is present as the [I-I-F]⁻ isomer, is a “non-classical” polyinterhalogen.

The propensity for halide anions to combine and form polyhalogen anions, such as I_3^- , has been known for around 200 years.¹ However, examples and spectroscopic characterisation of polyinterhalogen molecules, particularly open-shell species, have proven to be elusive.² To date, most spectroscopic characterisations of polyinterhalogen anions are as crystallised solids or have been performed in matrix isolation with a counter cation.^{2–4} There are no known vibrationally-resolved spectroscopic determinations of radical (open-shell) polyinterhalogen molecules. On the other hand, closed-shell polyinterhalogen molecules such as IF_3 , IF_5 , IF_7 and I_2F_4 are relatively stable compounds. Polyinterhalogen anions and their corresponding radical neutrals are appealing targets for gas-phase spectroscopy due to being textbook examples for valence-shell electron pair repulsion (VSEPR) theory and the Rundle-Pimentel scheme for hypervalency.^{5,6} They are also desirable systems to benchmark high-level quantum chemical calculations due to challenges associated with describing heavy atoms.^{7,8}

This paper reports a combined photoelectron spectroscopy and electronic structure theory study on the IF_2^- , I_2F^- , IF_2 and I_2F species. There have been several reports of synthesis, crystallography and spectroscopic characterisation of IF_2^- embedded in a

solid state matrix,^{9,10} and computational studies on IF_2^- and I_2F^- and related species,^{7,10–14} however, a recent review by Riedel and co-workers² noted “Thus far, ... $[\text{I}_2\text{F}]^-$ have not been detected experimentally.” While this statement might be true for condensed phases, I_2F^- appears to be a common product formed in hard Ar- CF_3I plasmas.

There are only a handful of examples of gas-phase spectroscopic studies on polyhalogen species, mostly targeting I_3^- . A series of pioneering investigations by Neumark and co-workers^{15–18} applied anion photoelectron spectroscopy with different laser sources to characterise vibronic properties of the radical neutral I_3 , and photodissociation dynamics of the parent anion. Several other studies have focussed on photodissociation dynamics.^{19–22} There have been two investigations on I_2Br^- probing the dissociation dynamics, with one of these studies providing a single-colour photoelectron spectrum at 267 nm.^{23,24} In the present work, we have coupled a jet-cooled plasma discharge source with time-of-flight mass spectrometry and anion photoelectron spectroscopy. This strategy allows for generation and mass selection of IF_2^- and I_2F^- , and determination of spectroscopic properties for the radical neutral IF_2 and I_2F species.

Intriguingly, radical polyinterhalogen molecules may have a significant bearing in industrial plasma processes. In particular, in the present study we produced the IF_2^- and I_2F^- species through plasmisation of an Ar- CF_3I mixture; this mixture and process is relevant in the semiconductor manufacture industry. In this industry, dry-etching plasma processes utilise the reaction between silicon wafers, which are the substrate for semiconductor and microprocessors, and fluorine atom and CF_3 radicals to etch the substrate at specific locations on the wafer.²⁵ Conventionally, these radical species are produced in a plasma consisting of an inert carrier gas seeded with CF_4 or C_3F_8 . In recent years, there has been growing interest in the use of CF_3I as an alternative to CF_4 because the use of CF_4 is restricted in many countries by the Montreal Protocol due to the molecule’s high global warming potential (GWP).^{26,27} For example, even though CF_3I is infrared active, CF_3I has an atmospheric residence life of ≈ 1 day

^a School of Chemistry, Norwich Research Park, University of East Anglia, Norwich NR4 7TJ, United Kingdom; E-mail: james.bull@uea.ac.uk

^b Department of Chemistry, Durham University, South Road, Durham DH1 3LE, United Kingdom

[†] Electronic Supplementary Information (ESI) available: Experimental methods, Theoretical methods, Calculated vertical excitation energies, Further Franck-Condon simulations. See DOI: 00.0000/00000000.

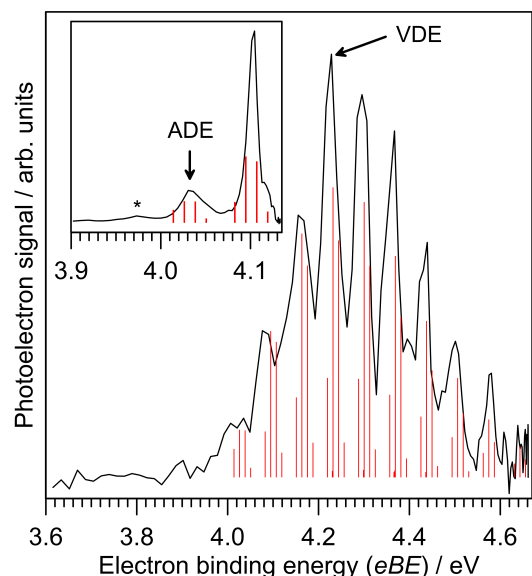


Fig. 1 Photoelectron spectra for IF_2^- at $h\nu = 4.66$ eV and 4.13 eV (inset) and Franck-Condon simulation (red sticks) of the $\text{D}_0 \leftarrow \text{S}_0$ detaching transition for the $[\text{F-I-F}]^-$ isomer. Electron binding energy ($e\text{BE}$) and electron kinetic energy ($e\text{KE}$) are related by $e\text{BE} = h\nu - e\text{KE}$, where $h\nu$ is the photon energy.

and corresponding $\text{GWP} \approx 1$.^{28,29} In contrast, CF_4 has a residence life of $\approx 50,000$ years and corresponding $\text{GWP} \approx 6,000$.^{29,30} Furthermore, although iodine atoms formed through UV photodissociation of CF_3I are exceptionally destructive towards stratospheric ozone,³¹ the short residence time of CF_3I released at terrestrial altitudes means that these molecules do not reach the stratosphere and contribute to ozone destruction. Ultimately, modelling and tuning of the etching process allows maximisation of process efficiency and minimisation of unwanted by-products. Theoretical plasma models require a detailed qualitative and quantitative understanding of the rich ion-molecule and electron-molecule chemistry that occurs in dry-etching plasmas (particularly the chemistry associated with fluorine-containing radicals).^{32–36}

IF_2^- & IF_2

Photoelectron spectra for IF_2^- are shown in Fig. 1, revealing a clear vibrational progression with spacing $600 \pm 50 \text{ cm}^{-1}$. The ADE, assigned as the lowest energy discernible detaching vibration in the $h\nu = 4.13$ eV spectrum, is $\text{ADE} = 4.03 \pm 0.02$ eV. The VDE, determined as the most intense vibration in the $h\nu = 4.66$ eV photoelectron spectrum, is $\text{VDE} = 4.23 \pm 0.04$ eV. These ADE and VDE values are most consistent with the calculated values (Table 1) for the $[\text{F-I-F}]^-$ isomer shown in Fig. 2; there was no evidence for the $[\text{F-F-I}]^-$ isomer, which was calculated to lie 2.74 eV (264 kJ mol^{-1}) higher in energy. Calculated ADE and VDE values for $[\text{F-I-F}]^-$ at the CCSD(T)//def2-TZVPD level of theory (Table 1) are both $\approx 4\%$ larger than the experimental values (similar to the result for I^- given in the ESI). Increasing the basis set to def2-QZVPD gave calculated values that were $\approx 6\%$ larger than the experimental values (Table 1). The better agreement with the smaller basis set is presumably a fortuitous cancellation of errors with some fraction of this associated with heavy atom/relativistic

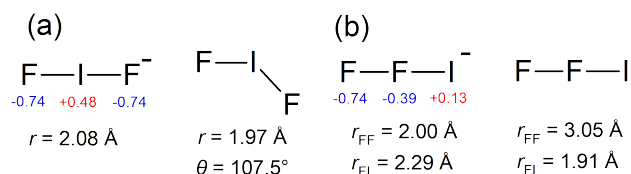


Fig. 2 Calculated equilibrium geometries at the CCSD(T)//def2-TZVPD level of theory for: (a) $[\text{F-I-F}]^-$ and $[\text{F-I-F}]$, and (b) $[\text{F-F-I}]^-$ and $[\text{F-F-I}]$. NBO charges (red and blue font) are given for the anions. The lowest energy isomer, $[\text{F-I-F}]^-$ has $\text{D}_{\infty h}$ symmetry, and $[\text{F-I-F}]$ has C_{2v} symmetry.

effects.

A Franck-Condon simulation of the photoelectron spectrum for the $[\text{F-I-F}]^-$ configuration is shown in Fig. 1 and has good agreement with the experimental spectrum. The simulation suggests that the vibrational progression is dominated by detaching transitions to the ν_2 (asymmetric stretch) and ν_3 (symmetric stretch) vibrations of the $[\text{F-I-F}]$ radical neutral species; calculated frequencies suggest that these two vibrations are nearly degenerate – see Table 2. We can therefore assign the experimental value of ν_2/ν_3 at $600 \pm 50 \text{ cm}^{-1}$ for the $[\text{F-I-F}]$ species. The small feature denoted by * in the inset in Fig. 1 is probably a hot band from hot ions produced in the plasma (see ESI). The ADE assignment should be reliable based on alignment of the VDE with the Franck-Condon simulation. It is worth noting that a Franck-Condon simulation of the photoelectron spectrum for the $[\text{F-F-I}]^-$ isomer (see vibrational frequencies for $[\text{F-F-I}]$ in Table 2) is inconsistent with the experimental spectra, lending support to the above assignment.

Calculated vertical excitation energies for the $[\text{F-I-F}]^-$ isomer are given in Table S1 in the ESI. These data show that although an excited state resonance is energetically accessible with a $h\nu = 4.66$ eV photon, oscillator strengths are zero and therefore the photoelectron spectra in Fig. 1 should be described by direct photodetachment to the ground electronic state of the neutral. Calculated vertical excitation energies for neutral $[\text{F-I-F}]$, i.e. electronic excited states of $[\text{F-I-F}]$ at the $[\text{F-I-F}]^-$ equilibrium geometry, suggested that only the ground electronic state is accessible using the photon energies in Fig. 1, consistent with a single vibrational progression.

I_2F^- & I_2F

Photoelectron spectra for I_2F^- at three photon energies are shown in Fig. 3a, revealing two detachment bands. The lower bind-

	$[\text{F-I-F}]^-$	$[\text{F-F-I}]^-$	$[\text{I-I-F}]^-$	$[\text{I-F-I}]^-$
ADE_e	4.03 ± 0.02	—	4.04 ± 0.06	—
VDE_e	4.23 ± 0.04	—	4.23 ± 0.04	—
ADE_c	4.14 (4.15)	3.17	4.22 (4.28)	3.19 (3.38)
VDE_c	4.40 (4.50)	3.88	4.40 (4.55)	3.96 (4.08)

Table 1 Experimental (subscript e) and calculated (subscript c) electron detachment energies for IF_2^- and I_2F^- . Calculations are at the CCSD(T)/def2-TZVPD level of theory (CCSD(T)/def2-QZVPD in parentheses). Experimental uncertainties account for calibration uncertainty from I^- and pixel-to-energy conversion in image reconstruction.

Mode	[F-I-F] ⁻	[F-I-F]	[F-F-I] ⁻	[F-F-I]
ν_1	193 ^a	96 ^a	137 ^a	18 ^a
ν_2	193 ^a	547 ^c	137 ^a	18 ^a
ν_3	401 ^c	548 ^b	230 ^c	38 ^b
ν_4	437 ^b	—	247 ^b	614 ^c

Table 2 Calculated harmonic vibrational frequencies for IF_2^- and IF_2 at the CCSD(T)//def2-TZVPD level of theory in units of cm^{-1} . ^aBend. ^bSymmetric stretch. ^cAsymmetric stretch. [F-F-I] has $C_{\infty h}$ symmetry. Note the symmetric vs asymmetric stretch mode ordering.

ing energy band has $\text{ADE}=4.04\pm0.06$ eV and $\text{VDE}=4.23\pm0.04$ eV. There is no clear vibrational structure. The higher binding energy band has $\text{ADE2}=4.60\pm0.04$ eV and $\text{VDE2}=4.68\pm0.04$ eV, and vibrational spacing of 550 ± 100 cm^{-1} .

The ADE and VDE parameters for the lower energy band are consistent with the calculated values for the $[\text{I-I-F}]^-$ isomer (Table 1), again with a small overestimation by theory. The two bands in Fig. 3a are thus assigned to detaching transitions to the ground and excited electronic states of the $[\text{I-I-F}]$ radical neutral. Equilibrium geometries for the I_2F^- and I_2F isomers are shown in Fig. 4; the $[\text{I-I-F}]^-$ isomer is more stable by 1.35 eV (130 kJ mol^{-1}), consistent with it being the predominant isomer formed in the plasma. There was no evidence for the $[\text{I-F-I}]^-$

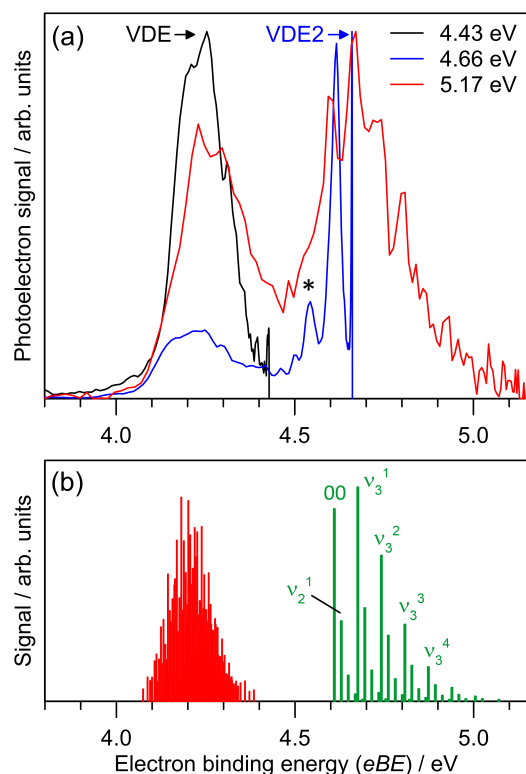


Fig. 3 (a) Photoelectron spectra for I_2F^- at $h\nu=4.43$, 4.66 and 5.17 eV. Electron binding energy ($e\text{BE}$) and electron kinetic energy ($e\text{KE}$) are related by $e\text{BE} = h\nu - e\text{KE}$, where $h\nu$ is the photon energy. (b) Franck-Condon simulation of direct photodetachment to the first neutral electronic states. The simulations suggest that the $\text{D}_0 \leftarrow \text{S}_0$ detaching transition is dominated by low frequency modes and combination bands. See ESI for Franck-Condon simulations at higher temperatures which assign the feature denoted by * to a hot band.

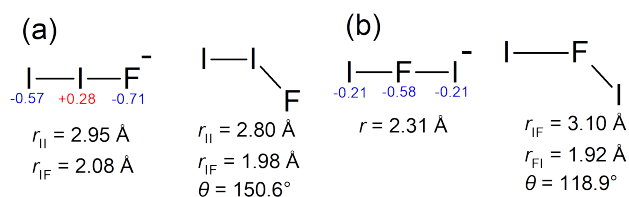


Fig. 4 Calculated equilibrium geometries at the CCSD(T)//def2-TZVPD level of theory for: (a) $[\text{I-I-F}]^-$ and $[\text{I-I-F}]$, and (b) $[\text{I-F-I}]^-$ and $[\text{I-F-I}]$. NBO charges (red and blue font) are given for the anions. The lowest energy isomer, $[\text{I-I-F}]^-$ has $C_{\infty h}$ symmetry, and $[\text{I-I-F}]$ has C_s symmetry.

isomer when photoelectron spectra were recorded at photon energies below the ADE for the $[\text{I-I-F}]^-$ isomer. A Franck-Condon simulation of the photoelectron spectrum to give the ground electronic state of the neutral (red in Fig. 3b) shows no clear vibrational structure, consistent with experimental spectrum.

The second, higher binding energy photodetachment band in Fig. 3a was assigned to direct photodetachment to the A_1 state of the $[\text{I-I-F}]$ species. Geometry optimisation of this state at the EOM-CCSD//def2-TZVPD level of theory produced a linear ($C_{\infty h}$ symmetry) structure with $r_{\text{II}}=3.02$ Å and $r_{\text{IF}}=1.96$ Å. Using this structure, ADE2 was calculated at 4.79 eV (0.57 eV difference to ADE), which is again $\approx 4\%$ larger than experiment. Calculated vibrational frequencies for the A_1 state state are $\nu_1=109$ cm^{-1} (bend), $\nu_2=157$ cm^{-1} (symmetric stretch) and $\nu_3=530$ cm^{-1} (asymmetric stretch). A Franck-Condon simulation of the photoelectron spectrum associated with formation of the A_1 state of the neutral is shown in green in Fig. 3b, and predicts that the vibrational structure is predominantly from the ν_3 mode. The feature denoted by * in Fig. 3a at $e\text{BE}=4.56$ eV is assigned to hot band signal (see ESI).

Calculated vertical excitation energies for the I_2F^- isomers are given in Table S1 in the ESI. For the $[\text{I-I-F}]^-$ isomer, the first few electronic states have very low oscillator strengths and the photoelectron spectra for photon energies < 5 eV should be dominated by direct photodetachment. There is, however, a bright A_1 -symmetry resonance with a calculated vertical excitation energy in the 5.1 – 5.2 eV range (see ESI). Fortunately, the photoelectron spectra for both of these species are at lower binding energies and should not interfere with the spectra shown in Fig. 3. It is worth noting that the issue of bright excited state resonances in the detachment continuum is more serious for I_3^{15-18} and larger polyinterhalogen anions due to an increased density of electronic states, hampering measurement of photoelectron spectra.

Polyinterhalogen molecules can be divided into two categories,

Mode	$[\text{I-I-F}]^-$	$[\text{I-I-F}]$	$[\text{I-F-I}]^-$	$[\text{I-F-I}]$
ν_1	124 ^a	102 ^a	99 ^a	20 ^a
ν_2	124 ^a	158 ^b	99 ^a	72 ^b
ν_3	136 ^b	502 ^c	120 ^b	606 ^c
ν_4	402 ^c	—	229 ^c	—

Table 3 Calculated harmonic vibrational frequencies for I_2F^- and I_2F at the CCSD(T)//def2-TZVPD level of theory in units of cm^{-1} . ^aBend. ^bSymmetric stretch. ^cAsymmetric stretch.

classical and non-classical.³ The former is defined as containing an electropositive central atom surrounded by electronegative halogen atoms, e.g. $[\text{F-I-F}]^-$. In contrast, for a non-classical polyinterhalogen molecule, the central halide is more electronegative than the coordinating dihalogen or interhalogen molecules, e.g. $\text{Cl}(\text{I}_2)_4^-$ in which four I_2 molecules coordinate a Cl^- .^{3,37} Interestingly, the $[\text{I-I-F}]^-$ species satisfies the non-classical criterion, possessing covalent-like bond lengths and a terminal F^- coordinated to an I_2 molecule (see NBO charges in Fig. 4). Similar arguments have been made about the $[\text{I-I-Cl}]^-$ species in the solid state.^{3,38} Finally, it is worth noting that Mabbs *et al.*²³ reported a 267 nm photoelectron spectrum of I_2Br^- , which presumably exists as the $[\text{I-I-Br}]^-$ isomer and also is an example of a non-classical polyinterhalogen, however, the photoelectron spectrum showed no vibrational structure.

Our experiments suggest that IF_2^- , I_2F^- and most likely the corresponding neutrals are amongst the major products formed in a hard $\text{Ar-CF}_3\text{I}$ plasma. In the semiconductor manufacture industry, radical neutrals including IF_2 and I_2F formed in the etching plasma may collide and react with etchant substrates. In this context, the bond dissociation energy (BDE) to produce a fluorine atom is related to etching efficiency – a lower BDE gives enhanced etching capacity. For the $[\text{F-I-F}]$ and $[\text{I-I-F}]$ isomers, BDEs for fluorine atoms were calculated at 1.74 eV and 1.35 eV, respectively, using the CCSD(T)//def2-TZVPD level of theory. For comparison, the calculated BDE for a fluorine atom from the CF_3 radical is 3.51 eV, which is substantially larger than that for the $[\text{F-I-F}]$ and $[\text{I-I-F}]$ isomers and suggests that polyinterhalogen radicals formed in hard $\text{Ar-CF}_3\text{I}$ plasmas are important etching species.

In summary, this paper has reported a combined photoelectron spectroscopy and electronic structure theory study on the IF_2^- , IF_2 , IF_2^- and I_2F species, determining the gas-phase structures and electron detachment parameters for the anions and vibrational structure of the neutrals. This determination provides the first vibrationally-resolved spectroscopic characterisation of isolated polyinterhalogen radicals. Whereas IF_2^- as the $[\text{F-I-F}]^-$ isomer is a classical polyinterhalogen molecule, IF_2^- as the $[\text{I-I-F}]^-$ isomer is classified as a non-classical polyinterhalogen molecule. Theoretical modelling of these species has shown that the CCSD(T)//def2-TZVPD level of theory, on average, predicts ADE and VDE parameters to within $\approx 4\%$ of experiment, however, this level of theory suffers from some degree of Pauling-point agreement. The IF_2 and I_2F radicals may play an important role in plasma etching processes due to having low bond dissociation energies to give fluorine atoms.

Funding was provided by a University of East Anglia start-up allowance (to JNB) and the European Research Council (Starting Grant 306536 to JRRV). Electronic structure calculations were carried out on the High Performance Computing Cluster supported by the Research and Specialist Computing Support service at the University of East Anglia. Prof. Dr. Sebastian Riedel, Freie Universität Berlin, is thanked for insightful discussions.

Conflicts of interest

There are no conflicts to declare.

Notes and references

- J. Pelletier and J. B. Caventou, *Ann. Chim. Phys.*, 1819, **10**, 142–176.
- K. Sonnenberg, L. Mann, F. A. Redeker, B. Schmidt and S. Riedel, *Angew. Chem. Int. Ed.*, 2020, doi:10.1002/anie.201903197.
- H. Haller and S. Riedel, *Z. Anorg. Allg. Chem.*, 2014, **640**, 1281–1291.
- P. H. Svensson and L. Kloos, *Chem. Rev.*, 2003, **103**, 1649–1684.
- R. J. Hach and R. E. Rundle, *J. Am. Chem. Soc.*, 1951, **73**, 4321–4324.
- G. C. Pimente, *J. Chem. Phys.*, 1951, **19**, 446–448.
- G. A. Landrum, N. Goldberg and R. Hoffmann, *J. Chem. Soc., Dalton Trans.*, 1997, 3605–3613.
- P. Pykkö, *Ann. Rev. Phys. Chem.*, 2012, **63**, 45–64.
- X. Zhang and K. Seppelt, *Z. Anorg. Allg. Chem.*, 1997, **623**, 491–500.
- K. O. Christe, W. W. Wilson, G. W. Drake, M. A. Petrie and J. A. Boatz, *J. Fluor. Chem.*, 1998, **88**, 185–189.
- Y. Ogawa, O. Takahashi and O. Kikuchi, *J. Mol. Struct. THEOCHEM*, 1998, **429**, 187–196.
- D. A. Dixon, D. J. Grant, K. O. Christe and K. A. Peterson, *Inorg. Chem.*, 2008, **47**, 5485–5494.
- K. S. Thanthiriatte, M. Vasilu, D. A. Dixon and K. O. Christe, *Inorg. Chem.*, 2012, **51**, 10966–10982.
- F. A. Redeker, A. Kropman, C. Müller, S. E. Zewge, H. Beckers, B. Paulus and S. Riedel, *J. Fluor. Chem.*, 2018, **216**, 81–88.
- T. R. Taylor, K. R. Asmis, M. T. Zanni and D. M. Neumark, *J. Chem. Phys.*, 1999, **110**, 7607–7609.
- M. T. Zanni, B. J. Greenblatt, A. V. Davis and D. M. Neumark, *J. Chem. Phys.*, 1999, **111**, 2991–3003.
- H. Choi, R. T. Bise, A. A. Hoops and D. M. Neumark, *J. Chem. Phys.*, 2000, **113**, 2255.
- H. Choi, T. R. Taylor, R. T. Bise, A. A. Hoops and D. M. Neumark, *J. Chem. Phys.*, 2000, **113**, 8608.
- L. Zhu, K. Takahashi, M. Saeki, T. Tsukuda and T. Nagata, *Chem. Phys. Lett.*, 2001, **350**, 223–239.
- A. A. Hoops, J. R. Gascooke, A. E. Faulhaber, K. E. Kautzman and D. M. Neumark, *J. Chem. Phys.*, 2004, **120**, 7901–7909.
- R. Nakanishi, N. Saitou, T. Ohno, S. Kowashi, S. Yabushita and T. Nagata, *J. Chem. Phys.*, 2007, **126**, 204311.
- J. A. Gibbard and R. E. Continetti, *Faraday Discuss.*, 2019, **217**, 203–219.
- R. Mabbs, K. Pichugin, E. Surber and A. Sanov, *J. Chem. Phys.*, 2004, **121**, 265.
- P. E. Crider, A. W. Harrison and D. M. Neumark, *J. Chem. Phys.*, 2011, **134**, 134306.
- J. W. Coburn and H. F. Winters, *J. Vac. Sci. Tech.*, 1979, **16**, 391–403.
- P. Forster, V. Ramaswamy, P. Artaxo, T. Bernsten, R. Betts, D. W. Fahey, J. Haywood, J. Lean, D. C. Lowe, G. Myhre, J. Nganga, R. Prinn, G. Raga, M. Schulz and R. Van Dorland, in *Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change*, ed. S. Solomon, D. Qin, M. Manning, Z. Chen, M. Marquis, K. Averyt, M. Tignor and H. Miller, Cambridge University Press, Cambridge, United Kingdom, 2007, ch. 2. Changes in Atmospheric Constituents and in Radiative Forcing, p. 141.
- Ø. Hodnebrog, M. Etminan, J. S. Fuglestad, G. Marston, G. Myhre, C. J. Nielsen, K. P. Shine and T. J. Wallington, *Rev. Geophys.*, 2013, **51**, 300–378.
- S. Solomon, J. B. Burckholder, A. R. Ravishankara and R. R. Garcia, *J. Geophys. Res.*, 1994, **99**, 20929.
- A. K. Jain, B. P. Briegleb, K. Minschwaner and D. J. Wuebbles, *J. Geophys. Res.: Atm.*, 2000, **105**, 20773–20790.
- C. M. Roehl, D. Boglu, C. BrÄijhl and G. K. Moortgat, *Geophys. Res. Lett.*, 1995, **22**, 815–818.
- S. Solomon, R. R. Garcia and A. R. Ravishankara, *J. Geophys. Res.*, 1994, **99**, 20491.
- L. G. Christophorou and J. K. Olthoff, *J. Phys. Chem. Ref. Data*, 2000, **29**, 553–569.
- J. N. Bull, M. Bart, C. Vallance and P. W. Harland, *Phys. Rev. A*, 2013, **88**, 062710.
- J. N. Bull, J. W. L. Lee and C. Vallance, *Phys. Rev. A*, 2017, **91**, 022704.
- G. Mensa-Bonsu, D. J. Tozer and J. R. R. Verlet, *Phys. Chem. Chem. Phys.*, 2018, **21**, 13977–13985.
- H. Köckert, D. Heathcote, J. W. L. Lee, W. Zhou, V. Richardson and C. Vallance, *Phys. Chem. Chem. Phys.*, 2019, **21**, 14296–14305.
- C. Walbaum, M. Richter, U. Sachs, I. Pantenburg, S. Riedel, A. Mudring and G. Meyer, *Angew. Chem. Int. Ed.*, 2013, **125**, 12965–12968.
- Y.-Q. Wang, Z.-M. Wang, C.-S. Liao and C.-H. Yan, *Act. Crystallog. Sect. C*, 1999, **55**, 1503–1506.